



TITLE:

# Gastrointestinal passage time of seeds ingested by captive Japanese martens *Martes melampus*

AUTHOR(S):

Tsuji, Yamato; Shiraishi, Toshiaki; Miura, Sayako

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1     Gastrointestinal passage time of seeds ingested by captive Japanese martens *Martes*

2     *melampus*

3     Yamato TSUJI, Toshiaki SHIRAISHI, and Sayako MIURA

4             The time it takes for ingested seeds to pass through the gut of animals is an important

5     aspect of endozoochorous seed dispersal because it influences seed dispersal distance.

6     Variations in the physical characteristics of seeds, such as their weight, volume, and

7     specific gravity, can affect their movement through the gastrointestinal system of a given

8     animal. We conducted feeding experiments with captive Japanese martens, *Martes*

9     *melampus* ( $n = 4$ ), at Toyama Municipal Family Park Zoo, central Japan, to examine the

10    effects of the physical characteristics of seeds on their passage times. The mean ( $\pm$  SD)

11    transit time, mean retention time, and time of last appearance of four different types of

12    commercial seeds were  $2.6 \pm 0.3$  hr (range: 0.6 – 5.4),  $9.7 \pm 1.1$  hr (3.8 – 17.3), and  $23.8$

13     $\pm 3.1$  hr (12.2 – 51.8), respectively. All of these values are greater than those found during

14    previous experiments conducted with mustelids. Similar to previous studies, however,

15    none of these passage time variables was correlated with the physical characteristics of

16 seeds. Our results thus indicate that martens disperse seeds of different plant species,  
17 whose size, volume, and specific gravity all fall within the range of those used in the  
18 present study, from parent plants at similar distances.  
19  
20 Primate Research Institute, Kyoto University, Aichi 484-8506, Japan, e-mail:  
21 ytsuji@pri.kyoto-u.ac.jp (YT); Toyama Municipal Family Park Zoo, Toyama 930-0151,  
22 Japan (TS, SM)  
23  
24 Key words: feeding experiments, *Martes melampus*, physical characteristics, seed  
25 dispersal  
26 Running page headline: Seed passage time of Japanese martens

## 27 Introduction

28       The physical characteristics of seeds, such as size, weight and specific gravity, vary  
29 among plant species. These variations can have a significant impact on passage time  
30 through the gut of a given animal (Traveset 1998). Leavy and Grajal (1991), for example,  
31 showed a negative correlation between the size of seeds ingested by cedar waxwings  
32 *Bombycilla cedrorum* and their subsequent passage times. Smaller seeds would thus be  
33 dispersed farther from parent plants. In contrast, Julliot (1996) provided evidence of a  
34 positive correlation between seed size and passage time in red howler monkeys *Alouatta*  
35 *seniculus*. To further confuse the issue, the relationship between passage time and  
36 specific gravity, another physical characteristic relevant to seed dispersal, was positive  
37 for seeds ingested by Japanese macaques *Macaca fuscata* (Tsuji *et al.* 2010) but negative  
38 for those ingested by two species of tamarin monkeys *Saguinus mystax* and *S. fuscicollis*  
39 (Garber 1986). Other studies have found no clear relationships between the physical  
40 characteristics of seeds and their passage times (emus *Dromaius novahollandiae*: Wilson  
41 1989; arctic foxes *Alopex lagopus*: Graae *et al.* 2004; Pampa foxes *Lycalopex*

42 *gymnocercus* and crab-eating foxes *Cerdocyon thous*: Varela and Bucher 2006). Thus, it  
43 appears that the effects of physical characteristics on seed passage time vary among  
44 animal species, and this may reflect variation in the size and/or morphology of the  
45 digestive system and the digestive strategies of the animals under investigation.

46 Mustelids are generally regarded as carnivorous, but they also commonly consume  
47 large amounts of fruit (Rosalino and Santos-Reis 2009), often voiding intact seeds with  
48 their feces (pine martens *Martes martes*: Schaumann and Heinken 2002; stone martens *M.*  
49 *foina*: Schaumann and Heinken 2002; Japanese martens *M. melampus*: Arai *et al.* 2002;  
50 Koike *et al.* 2008; yellow-throated martens *M. flavigula*: Zhou *et al.* 2008a). The passage  
51 of seeds through the guts of martens can have a positive effect on germination and/or  
52 seedling growth (Hickey *et al.* 1999, Schaumann and Heinken 2002, but see Rosalino *et*  
53 *al.* 2010). Therefore, the marten appears to be an important seed disperser within its  
54 habitat (Hickey *et al.* 1999). Since mustelids travel between several hundred meters and  
55 several kilometers daily (Hickey *et al.* 1999, Zalewski *et al.* 2004), and have relatively  
56 large home ranges amounting to between 1 and 4 km<sup>2</sup> (Zalewski and Jedrzejewski 2006),

57 variation in passage time can markedly affect seed dispersal distance, which in turn  
58 affects the seed shadow of a given plant species. Thus, evaluating the effects of the  
59 physical characteristics of seeds on passage time through marten digestive tracts is  
60 essential to understanding their role in endozoochorous seed dispersal.

61 In this study we conducted feeding experiments with captive Japanese martens,  
62 which are considered an important mammalian seed disperser throughout Japanese  
63 forests (Kusui and Kusui 1998, Otani 2002, Koike *et al.* 2008, Tsuji *et al.* in press). We  
64 tested one simple prediction: that seed passage time through the gut of Japanese martens  
65 differs among seed types because of variation in their weights, volumes, and specific  
66 gravities.

67

## 68 Materials and methods

69 Four adult martens (three males and one female) with which we conducted our  
70 feeding experiments were housed at the Toyama Municipal Family Park Zoo, central  
71 Japan. All animals were reared in individual wire mesh pens (length 1.80 m × width 1.8 m

72 × height 2.8 m), and were active and in good condition. The estimated ages and body  
73 weights of the three animals for which data were available (two males and one female,  
74 respectively) were 16, 3, and 3 yr and 2.0, 1.8, and 1.0 kg. The female marten was neither  
75 lactating nor pregnant during the study period. The martens were normally fed dead  
76 chicks, chicken heads, boiled sweet potatoes and fresh fruit (fresh weight: 230 – 300 g·  
77 individual<sup>-1</sup>) once a day (at 10:00). We only entered the pen during the experiments, to  
78 minimize stressing the animals and thereby affecting gut passage times. About one month  
79 before conducting the experiments, we set security cameras on the roofs of each pen to  
80 continuously monitor the movements of each animal. Videos were continuously recorded  
81 onto hard disks, so we could accurately record their defecation times and locations (see  
82 below).

83 In light of a previous review of fruit feeding by martens (Koike and Masaki 2008),  
84 and the results of our own recent study (Tsuji *et al.* in press), we selected the seeds of four  
85 commercial plant species (spinach, water spinach, kiwi fruits, and okra), with seed  
86 lengths ranging from 2.2 – 5.9 mm (Table 1) and shapes similar to those of many of the

87 plant species on which martens are known to ingest in the wild. Because of restrictions  
88 relating to experimentation with captive animals, we could not provide seeds collected in  
89 wild in this study. We measured the length, width, and height of 30 randomly-picked  
90 seeds with a vernier caliper (THS-30, Niigata Seiki Co., Japan) to the nearest 0.05 mm.  
91 We also weighed 30 randomly-picked dry seeds with an electric balance (UX4200H,  
92 Shimadzu Co., Japan) to the nearest 1 mg. We then calculated seed volume according to  
93 the formula:

94 
$$V = \pi R^2 \left( L - \frac{2}{3} R \right),$$

95 where  $R = (\text{seed width} + \text{height}) / 4$ , and  $L = \text{seed length}$  (Garber 1986). Finally, we  
96 calculated the specific gravity of seeds ( $\text{mg} \cdot \text{mm}^{-3}$ ) by dividing the seed volume by the  
97 seed weight (Table 1). After taking these measurements, we put seeds into the stomachs  
98 of dead chicks, since preliminary experiments showed that the martens would not ingest  
99 the seeds alone. We must thus accept the possibility that this treatment might have  
100 affected the results of our experiments. Information concerning the size, weight, and  
101 amount of seeds used in each experiment is shown in Table 1.



102        We conducted four experimental sessions between late September and early October,  
103        2010. We did not change the martens' housing conditions or diet composition during the  
104        experiments. Our methodology adhered to Japan's legal requirements for animal welfare.  
105        On the first day of a given experiment, each animal was fed as usual at 10:00, and we  
106        regularly entered the pen twice a day (10:00 and 16:00) to collect both non-ingested seeds  
107        and all feces found within the enclosure. We recorded the locations of defecation and  
108        cleaned the floor of the pen in order to facilitate subsequent monitoring of defecation  
109        events. We rinsed fecal samples with water immediately after collection and screened  
110        them using a 0.5 mm sieve, recording the number of intact seeds, defined as those with  
111        undamaged embryos, whenever present. We considered seeds that were ingested but not  
112        observed in faecal samples to be destroyed through mastication or digestion. We initiated  
113        subsequent experimental sessions after 24 hr had passed since the onset of the previous  
114        experimental session (i.e., 10:00). We recorded the time of defecation for each fecal  
115        sample from the videos taken for each animal.

116        According to previous studies of seed dispersal, the quantitative evaluation of seed

passage time is based on three parameters: transit time, mean retention time, and time of

last appearance. We obtained the mean retention time via the following formula:

$$MRT = \frac{\sum_{i=1} m_i t_i}{\sum_{i=1} m_i}$$

where  $MRT$  = mean retention time,  $m_i$  = the number of seeds excreted in the  $i$ th defecation

at time  $t_i$  (hr) after ingestion (Lambert 2002).

We tested the effects of seed type on the percentage of seeds recovered and on the

three passage time variables (transit time, mean retention time, and time of last

appearance), and the effects of individual marten on the passage times. We employed

Friedman's two-way ANOVAs to account for the non-normality of our data. We used

Spearman's correlation analyses to test for relationships between the physical

characteristics of seeds and both the percentage of seeds recovered and the three passage

time variables. We could not test for differences in seed recovery and passage time

variables between sexes because of our small sample size of four animals. We set the

level of significance for these analyses ( $\alpha$ ) at 0.05. All data analyses were conducted

using the statistical software R version 2.9.1 (R Development Core Team 2009).

## Results

The martens in our study readily and immediately consumed the dead chicks when presented with them, but not all of the seeds concealed therein were ingested because several dropped out of the chick during feeding (Table 1). The martens defecated more than half of their ingested seeds, The mean percentage of seeds recovered for each seed type ranged from 70 – 86 %, and did not vary significantly among seed types (Friedman's two-way ANOVA,  $\chi^2 = 4.3$ ,  $p = 0.233$ ,  $df = 3$ ) or among animals ( $\chi^2 = 4.9$ ,  $p = 0.181$ ,  $df = 3$ ). Furthermore, none of the physical dimensions of the seeds correlated with the percentage of seeds recovered (Spearman's correlation analyses, dry weight:  $r_s = -0.20$ ,  $p = 0.917$ ; volume:  $r_s = 0.40$ ,  $p = 0.750$ ; specific gravity:  $r_s = -0.40$ ,  $p = 0.750$ ,  $df = 2$  for all analyses).

The mean transit time, mean retention time, and time of last appearance were  $2.6 \pm 0.3$  hr (range: 0.6 – 5.4 hr, Fig. 1a),  $9.7 \pm 1.1$  hr (3.8 – 17.3 hr, Fig. 1b) and  $23.8 \pm 3.1$  hr (12.2 – 51.8 hr, Fig. 1c), respectively. We treated the three passage time variables as

independent of each other because there were no significant correlations among them (Spearman's correlation analyses, transit time vs mean retention time:  $r_s = -0.20$ ,  $p = 0.917$ ; transit time vs time of last appearance:  $r_s = -0.20$ ,  $p = 0.917$ ; mean retention time vs time of last appearance:  $r_s = -0.20$ ,  $p = 0.917$ ,  $df = 2$  for all analyses). None of these passage time variables differed significantly among seed types (Friedman's two-way ANOVA, transit time:  $\chi^2 = 1.5$ ,  $p = 0.682$ ; mean retention time:  $\chi^2 = 1.2$ ,  $p = 0.753$ ; time of last appearance:  $\chi^2 = 2.1$ ,  $p = 0.552$ ,  $df = 3$  for all analyses). Among these variables, mean retention time differed significantly between individual animals (Friedman's two-way ANOVA,  $\chi^2 = 9.9$ ,  $p = 0.019$ ,  $df = 3$ ), being longest in the heavier males and shortest in the lighter female. Finally, transit time and time of last appearance did not differ between individual animals (transit time:  $\chi^2 = 4.4$ ,  $p = 0.219$ ; time of last appearance:  $\chi^2 = 5.7$ ,  $p = 0.127$ ).

## Discussion

In this study, none of the three passage time variables differed significantly among

162 seed types in our experiments. Our results differ from those concerning other animal  
163 species such as birds (Leavy and Grajal 1991) and primates (Garber 1986, Tsuji *et al.*  
164 2010), in which the physical characteristics of seeds were shown to affect passage times.  
165 Our results imply that plant seeds with physical characteristics falling within the ranges  
166 of those used in the present study have the same probability of being dispersed by  
167 Japanese martens a similar distance away from parent plants. Previous studies of  
168 carnivorous mammals have also showed that the passage times of seeds ingested by their  
169 subject animals did not vary across seed types (Graae *et al.* 2004, Varela and Bucher 2006,  
170 Zhou *et al.* 2008b). Thus, the lack of a relationship between seed type and passage time  
171 appears to be a common trait of carnivorous mammals. However, we cannot deny the  
172 possibility that our results might have been affected by our small number of experiments.  
173 Another possibility is that factors that we did not consider in this study, such as fruit pulp  
174 properties (e.g. texture and chemistry), the amount of other fiber-rich compounds of the  
175 wild marten diet (e.g., meat and fur), and the amount and/or shape of seeds, may have  
176 masked the effects of seed size, volume, and specific gravity. Thus, further and more

detailed experimentation considering these effects should be conducted in the future (Traveset 1998). We also encourage cooperation between researchers and zoological gardens, as this will be invaluable in furthering our understanding of endozoochorous seed dispersal among carnivorous mammals.

By combining data on passage time and animal movement obtained through radio telemetry, Hickey *et al.* (1999) estimated a range of seed dispersal distances (i.e., seed shadow) created by American martens. However, there have been few quantitative studies to date evaluating ranging patterns among Japanese martens (Kawauchi *et al.* 2003, Okumura and Kitahara 2006). Furthermore, few studies have tested the effects of ingestion and passage through the digestive systems of Japanese martens on the germination and/or growth of those plant species. Given this lack of empirical data, our study can contribute to an estimation of the seed shadows created by Japanese martens. To be an effective seed disperser, fruit consumers cannot destroy the seeds they ingest, but they are also expected to enhance (or at least have a neutral effect on) seed germination during transit through the gut (Pollux *et al.* 2007, Rosalino *et al.* 2010). In

the future, testing the relationship between the seed dispersal distribution and seed performance (e.g. germination and seedling growth) in the field is a prerequisite to any evaluation of the efficacy of Japanese martens as seed dispersers.

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## 277    Figure legends

278    Figure 1: Relationships between seed type (n=4) and passage time variables: a) transit  
279    time, b) mean retention time, and c) time of last appearance of a seed. Filled circles  
280    indicate mean values and bars indicate standard deviations.

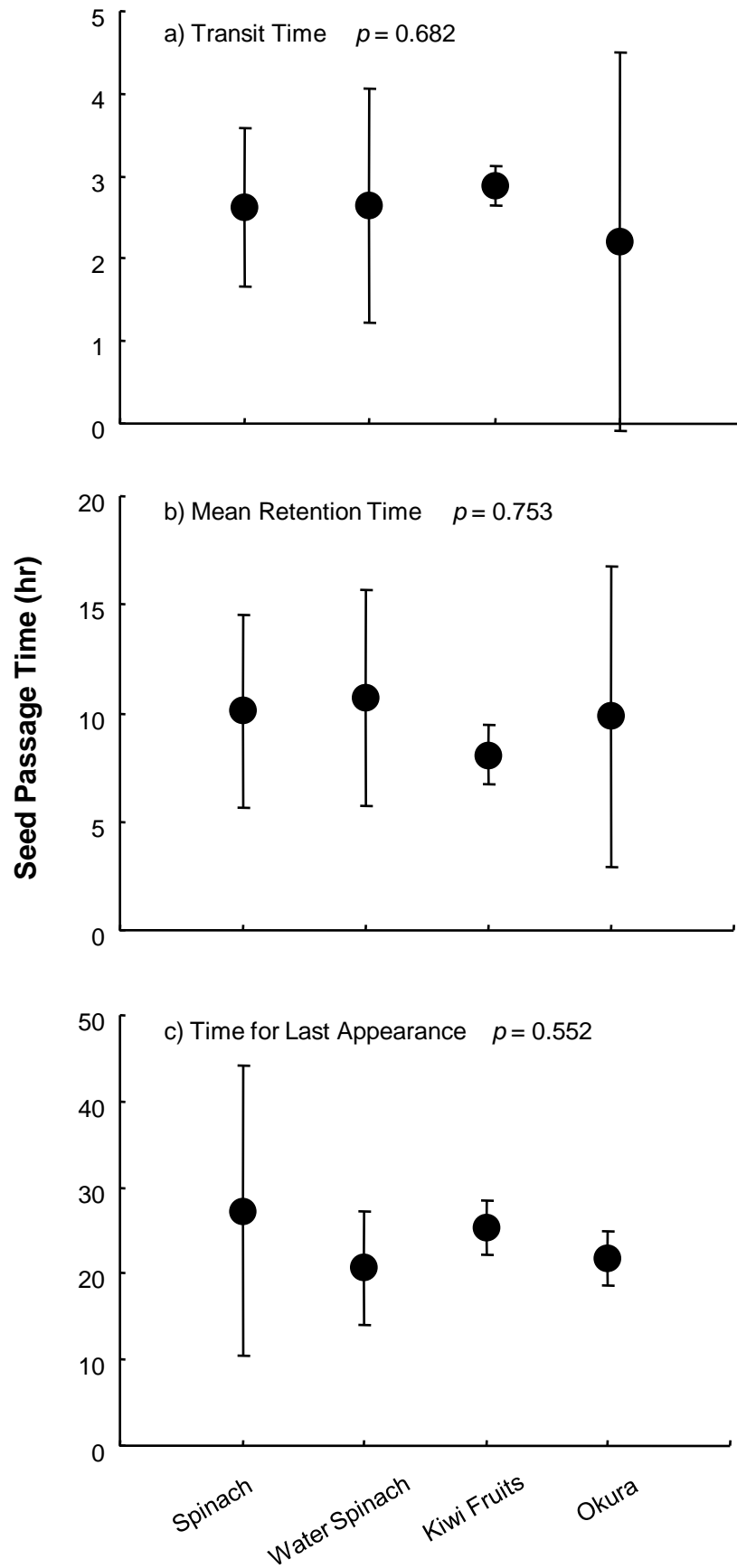


Table 1. The physical characteristics of commersial seeds used in feeding experiments and their fate.

Seed type	Physical characteristics of seeds							Dose	Seeds fate						
	Size (mm, mean ± SD)			Dry weight	Volume <sup>a)</sup>	Specific gravity <sup>b)</sup>	# seeds in a chick	Ingested		Defecated		% Recovery			
	Length	Width	Height	(mg, mean ± SD)	(mm <sup>3</sup> )	(mg·mm <sup>-3</sup> )		mean ± SD	range	mean ± SD	range	mean ± SD	range		
Spinach	3.58 ± 0.41	3.12 ± 0.33	2.26 ± 0.22	11.47 ± 3.17	15.25	0.75	40	39.8 ± 0.5	39 - 40	31.5 ± 3.5	28 - 35	79.2 ± 8.4	70.0 - 87.5		
Water spinach	5.92 ± 0.32	4.09 ± 0.34	3.55 ± 0.24	51.67 ± 8.53	53.29	0.97	40	39.5 ± 0.0	39 - 40	27.8 ± 4.0	23 - 32	70.4 ± 11.2	57.5 - 82.1		
Kiwi fruits	2.24 ± 0.21	1.23 ± 0.15	0.71 ± 0.08	1.00 ± 0.31	1.43	0.70	400	400.0 ± 0.0	-	278.8 ± 61.2	210 - 350	78.4 ± 10.0	52.5 - 87.8		
Okura	4.84 ± 0.31	4.23 ± 0.26	4.29 ± 0.31	43.69 ± 9.64	48.64	0.90	20	20.0 ± 0.0	-	16.3 ± 1.2	15 - 20	86.3 ± 10.3	75.0 - 100.0		

The sample number of seed measurements was 30 for each species.

a) Seed volume was calculated by using the following formula:  $V = \pi R^2 (L - 2/3R)$ ; V = volume, R = (width + height) / 4, L = length.

b) Calculated as dry weight / volume.